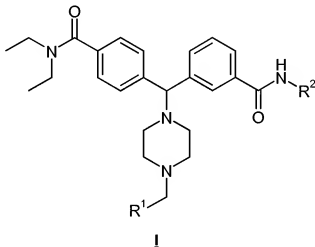


**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Currently amended) A compound of formula I, pharmaceutically acceptable salts thereof, or mixtures thereof:



wherein

R<sup>1</sup> is an C<sub>6-14</sub> aryl, heteroaryl, substituted aryl or substituted C<sub>3-20</sub> heteroaryl, wherein said aryl and heteroaryl are each independently and optionally substituted with one or more groups selected from C<sub>1-6</sub>hydrocarbon, -NO<sub>2</sub>, -OR, -Cl, -Br, -I, -F, -CF<sub>3</sub>, -C(=O)R, -C(=O)OH, -NH<sub>2</sub>, -SH, -NHR, -NR<sub>2</sub>, -SR, -SO<sub>3</sub>H, -SO<sub>2</sub>R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR<sub>2</sub>, -NRC(=O)R, oxo (=O), imino (=NR), thio (=S), and oximino (=N-OR), wherein each R is a C<sub>1-6</sub>hydrocarbonyl; and R<sup>2</sup> is hydrogen, optionally substituted C<sub>1-12</sub>alkyl, optionally substituted C<sub>6-12</sub>aryl, or optionally substituted C<sub>2-12</sub>heterocyclyl, wherein said alkyl, aryl, and heterocyclyl are each independently and optionally substituted with one or more groups selected from C<sub>1-6</sub>hydrocarbon, -NO<sub>2</sub>, -OR, -Cl, -Br, -I, -F, -CF<sub>3</sub>, -C(=O)R, -C(=O)OH, -NH<sub>2</sub>, -SH, -NHR, -NR<sub>2</sub>, -SR, -SO<sub>3</sub>H, -SO<sub>2</sub>R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR<sub>2</sub>, -NRC(=O)R, oxo (=O), imino (=NR), thio (=S), and oximino (=N-OR), wherein each R is a C<sub>1-6</sub>hydrocarbonyl.

2. (Original) A compound according to claim 1,

wherein R<sup>1</sup> is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl; triazolyl; pyrrolyl; thiazolyl; and N-oxido-pyridyl, optionally substituted with one or more groups selected from C<sub>1-6</sub>alkyl, halogenated C<sub>1-6</sub>alkyl, -NO<sub>2</sub>, -CF<sub>3</sub>, C<sub>1-6</sub> alkoxy, chloro, fluoro, bromo, and iodo; and

R<sup>2</sup> is hydrogen or methyl.

3. (Original) A compound according to claim 1, wherein R<sup>1</sup> is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; and thiazolyl, optionally substituted with one or more groups selected from C<sub>1-6</sub>alkyl, halogenated C<sub>1-6</sub>alkyl, -NO<sub>2</sub>, -CF<sub>3</sub>, C<sub>1-6</sub> alkoxy, chloro, fluoro, bromo, and iodo; and

R<sup>2</sup> is hydrogen or methyl.

4. (Original) A compound according to claim 1, wherein R<sup>1</sup> is selected from phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; and thiazolyl; and R<sup>2</sup> is hydrogen or methyl.

5. (Original) A compound according to claim 1, wherein the compound is selected from:

3-[(4-[(diethylamino)carbonyl]phenyl)(4-benzyl-piperazin-1-yl)methyl]benzamide;

3-[(4-[(diethylamino)carbonyl]phenyl)[4-(2-furylmethyl)-piperazin-1-yl)methyl]benzamide;

3-[(4-[(diethylamino)carbonyl]phenyl)[4-(phenylmethyl)-1-piperazinyl)methyl]-N-methylbenzamide; enantiomers thereof; and pharmaceutically acceptable salts thereof.

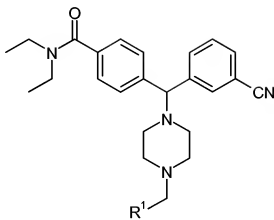
- 6-7. (Cancelled)

8. (previously presented) A pharmaceutical composition comprising a compound according to claim 1 and a pharmaceutically acceptable carrier.

9. (currently amended) A method for the therapy of pain in a warm-blooded animal, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1.

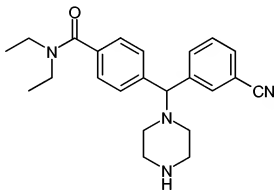
10. (cancelled)

11. (Currently Amended) A process for preparing a compound of formula II,



II

comprising of the step of reacting a compound of formula III:

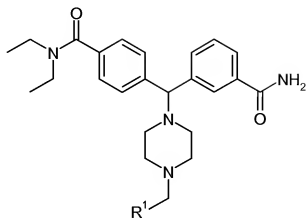


III

with R<sup>1</sup>-CHO to form the compound of formula II  
 wherein

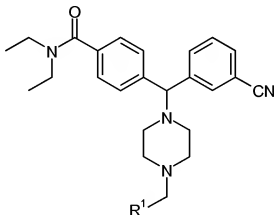
R<sup>1</sup> is an C<sub>5-14</sub> aryl, heteroaryl, substituted aryl or substituted C<sub>3-20</sub> heteroaryl, wherein said aryl and heteroaryl are each independently and optionally substituted with one or more groups selected from C<sub>1-6</sub> hydrocarbon, -NO<sub>2</sub>, -OR, -Cl, -Br, -I, -F, -CF<sub>3</sub>, -C(=O)R, -C(=O)OH, -NH<sub>2</sub>, -SH, -NHR, -NR<sub>2</sub>, -SR, -SO<sub>3</sub>H, -SO<sub>2</sub>R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR<sub>2</sub>, -NRC(=O)R, oxo (=O), imino (=NR), thio (=S), and oximino (=N-OR), wherein each R is a C<sub>1-6</sub> hydrocarbonyl.

12. (Currently Amended) A process for preparing a compound of formula IV,



**IV**

comprising: reacting a compound of formula II,



**II**

with an alkali metal hydroxide in non-aqueous solvent to form the compound of formula IV:  
 wherein

R<sup>1</sup> is an C<sub>6-14</sub> aryl, heteroaryl, substituted aryl or substituted C<sub>3-20</sub> heteroaryl, wherein said aryl and heteroaryl are each independently and optionally substituted with one or more groups selected from C<sub>1-6</sub> hydrocarbon, -NO<sub>2</sub>, -OR, -Cl, -Br, -I, -F, -CF<sub>3</sub>, -C(=O)R, -C(=O)OH, -NH<sub>2</sub>, -SH, -NHR, -NR<sub>2</sub>, -SR, -SO<sub>3</sub>H, -SO<sub>2</sub>R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR<sub>2</sub>, -NRC(=O)R, oxo (=O), imino (=NR), thio (=S), and oximino (=N-OR), wherein each R is a C<sub>1-6</sub> hydrocarbyl.